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        Apr 09
NEWS 4
        Apr 09
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                US Patent Applications available in IFICDB, IFIPAT, and IFIUDB
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                Records from IP.com available in CAPLUS, HCAPLUS, and ZCAPLUS
NEWS 7
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        Jun 03
                New e-mail delivery for search results now available
NEWS 10
        Jun 10 MEDLINE Reload
NEWS 11
        Jun 10 PCTFULL has been reloaded
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        Jul 02
                FOREGE no longer contains STANDARDS file segment
NEWS 13
        Jul 22 USAN to be reloaded July 28, 2002;
                 saved answer sets no longer valid
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        Jul 29
                Enhanced polymer searching in REGISTRY
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                NETFIRST to be removed from STN
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                CANCERLIT reload
        Aug 08
NEWS 17
        Aug 08
                PHARMAMarketLetter(PHARMAML) - new on STN
NEWS 18
        Aug 08 NTIS has been reloaded and enhanced
NEWS 19
        Aug 19
                Aquatic Toxicity Information Retrieval (AQUIRE)
                 now available on STN
NEWS 20
        Aug 19
                 IFIPAT, IFICDB, and IFIUDB have been reloaded
NEWS 21
                The MEDLINE file segment of TOXCENTER has been reloaded
        Aug 19
NEWS 22
        Aug 26
                Sequence searching in REGISTRY enhanced
NEWS 23
        Sep 03
                JAPIO has been reloaded and enhanced
NEWS 24
        Sep 16
                Experimental properties added to the REGISTRY file
NEWS 25
        Sep 16
                Indexing added to some pre-1967 records in CA/CAPLUS
NEWS 26
        Sep 16
                CA Section Thesaurus available in CAPLUS and CA
NEWS 27
        Oct 01
                CASREACT Enriched with Reactions from 1907 to 1985
NEWS 28
        Oct 21
                EVENTLINE has been reloaded
NEWS 29
        Oct 24
                BEILSTEIN adds new search fields
NEWS 30
        Oct 24
                Nutraceuticals International (NUTRACEUT) now available on STN
NEWS 31 Oct 25
                MEDLINE SDI run of October 8, 2002
NEWS EXPRESS
             October 14 CURRENT WINDOWS VERSION IS V6.01,
             CURRENT MACINTOSH VERSION IS V6.0a(ENG) AND V6.0Ja(JP),
             AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002
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STRUCTURE FILE UPDATES: 29 OCT 2002 HIGHEST RN 467418-81-1 DICTIONARY FILE UPDATES: 29 OCT 2002 HIGHEST RN 467418-81-1

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=> s 162359-55-9/rn or 162359-55-9/crn

1 162359-55-9/RN

1 162359-55-9/CRN

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=> s l1
'RN' IS NOT A VALID FIELD CODE
'CRN' IS NOT A VALID FIELD CODE
           632 L1
=> s FTY 720
           366 FTY 720
=> s 12 or 13
           643 L2 OR L3
=> s immunostimul? or immunoenhanc?
         48360 IMMUNOSTIMUL? OR IMMUNOENHANC?
=> s 14 and 15
             1 L4 AND L5
=> d
     ANSWER 1 OF 1 EMBASE COPYRIGHT 2002 ELSEVIER SCI. B.V.
L6
     2002060333 EMBASE
AN
TΤ
     L-selectin-dependent lymphoid occupancy is required to induce
     alloantigen-specific tolerance.
     Bai Y.; Liu J.; Wang Y.; Honig S.; Qin L.; Boros P.; Bromberg J.S.
AU
     Dr. J.S. Bromberg, Mount Sinai School of Medicine, Box 1104, One Gustave
CS
     L. Levy Place, New York, NY 10029-6574, United States.
     jon.bromberg@mountsinai.org
SO
     Journal of Immunology, (15 Feb 2002) 168/4 (1579-1589).
     Refs: 48
     ISSN: 0022-1767 CODEN: JOIMA3
CY
     United States
DT
     Journal; Article
FS
     018
             Cardiovascular Diseases and Cardiovascular Surgery
     026
             Immunology, Serology and Transplantation
     030
             Pharmacology
             Drug Literature Index
     037
LA
     English
SL
     English
=> s viral or antiviral or antiviru? or viru?
   3 FILES SEARCHED...
       2026272 VIRAL OR ANTIVIRAL OR ANTIVIRU? OR VIRU?
=> s 17 and 14
            27 L7 AND L4
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PROCESSING COMPLETED FOR L8
             21 DUP REM L8 (6 DUPLICATES REMOVED)
=> d ibib abs 16-21
     ANSWER 16 OF 21
                         MEDLINE
                                                         DUPLICATE 3
ACCESSION NUMBER:
                    2000281664
                                   MEDLINE
DOCUMENT NUMBER:
                    20281664
                              PubMed ID: 10820254
TITLE:
                    FTY720 immunosuppression impairs effector T cell peripheral
                    homing without affecting induction, expansion, and memory.
AUTHOR:
                    Pinschewer D D; Ochsenbein A F; Odermatt B; Brinkmann V;
                    Hengartner H; Zinkernagel R M
```

Institute of Experimental Immunology and Laboratory for

CORPORATE SOURCE:

Special Techniques, Department of Pathology, University

Hospital, Zurich, Switzerland.

SOURCE: JOURNAL OF IMMUNOLOGY, (2000 Jun 1) 164 (11) 5761-70.

Journal code: 2985117R. ISSN: 0022-1767.

PUB. COUNTRY: United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Abridged Index Medicus Journals; Priority Journals

ENTRY MONTH: 200006

ENTRY DATE: Entered STN: 20000629

Last Updated on STN: 20000629 Entered Medline: 20000621

AB FTY720 (2-amino-2-(2-[4-octylphenyl]ethyl)-1,3-propanediol hydrochloride) prolongs survival of solid organ allografts in animal models. Mechanisms of FTY720 immunomodulation were studied in mice infected with lymphocytic choriomeningitis virus (LCMV) to assess T cell responses or with vesicular stomatitis virus to evaluate Ab responses. Oral FTY720 (0.3 mg/kg/day) did not affect LCMV replication and specific CTL and B cells were induced and expanded normally. Moreover, the anti-viral humoral immune responses were normal. However, FTY720 treatment showed first a shift of overall distribution of CTL from the spleen to peripheral lymph nodes and lymphocytopenia was observed. This effect was reversible within 7-21 days. Together with unimpaired T and B cell memory after FTY720 treatment, this finding rendered enhancement of lymphocyte apoptosis by FTY720 in vivo unlikely. Secondly, the delayed-type hypersensitivity reaction to a viral MHC class I-presented peptide was markedly reduced by FTY720. These results were supported by impaired circulation of LCMV specific TCR transgenic effector lymphocytes in the peripheral blood and reduced numbers of tissue infiltrating CTL in response to delayed-type hypersensitivity reaction. Thirdly, in a CD8+ T cell-mediated diabetes model in a transgenic mouse expressing the LCMV glycoprotein in the islets of the pancreas, FTY720 delayed or prevented disease by reducing islet-infiltrating CTL. Thus, FTY720 effectively reduced recirculation of CD8+ effector T cells and their recruitment to peripheral lesions without affecting the induction and expansion of immune responses in secondary lymphoid organs. These properties may offer the potential to treat ongoing organ-specific T cell-mediated immunopathologic disease.

L9 ANSWER 17 OF 21 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 2000:896183 CAPLUS

DOCUMENT NUMBER: 135:55693

TITLE: Perioperative administration of FTY720 and CTLA4IG in

rat heart transplantation

AUTHOR(S): Ohba, M.; Li, X.-K.; Kita, Y.; Tamura, A.; Enosawa,

S.; Sasakuri, S.; Ogoshi, S.; Amemiya, H.; Suzuki, S. Department of Experimental Surgery and Bioengineering,

National Children's Medical Research Center, Tokyo,

Japan

SOURCE: Transplantation Proceedings (2000), 32(7), 2024-2025

CODEN: TRPPA8; ISSN: 0041-1345

PUBLISHER: Elsevier Science Inc.

DOCUMENT TYPE: Journal LANGUAGE: English

CORPORATE SOURCE:

AB A study was conducted to examine the in vitro proliferation activity of lymphocytes from recipients transfected with adenovirus vectors contg. CTLA4Ig-gene (AdCTLA4Ig) and FTY720 administered in a rat model of allogeneic heart transplantation. The administration of FTY720 or AdCTLA4Ig resulted in significant prolongation of allograft survival. The combination therapy with FTY720 and AdCTLA4Ig caused further prolongation effects on graft survival time. The in vitro proliferation activity of lymphocytes to donor cells were completely inhibited early after grafting

in both FTY720-treated recipients and AdCTLA4Ig-treated ones. FTY720-treated recipients showed a marked suppression in lymphocyte response 14 days after grafting, whereas the lymphocytes from AdCTLA4Iq-treated recipients recovered the response despite absence of a rejection episode. In addn., a remarkable inhibition of mixed lymphocyte reaction was obsd. in the lymphocytes from recipients with combination therapy.

REFERENCE COUNT:

THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 18 OF 21 USPATFULL

ACCESSION NUMBER:

1999:166606 USPATFULL

TITLE:

Compositions and methods of using compositions with

accelerated lymphocyte homing immunosuppressive

properties

INVENTOR(S):

Chiba, Kenji, Fukuoka, Japan Adachi, Kunitomo, Fukuoka, Japan

PATENT ASSIGNEE(S):

Yoshitomi Pharmaceutical Industries, Ltd., Osaka, Japan

(non-U.S. corporation)

NUMBER KIND DATE -----US 6004565 19991221

PATENT INFORMATION: APPLICATION INFO.:

US 1997-933738

19970923 (8)

NUMBER DATE -----

PRIORITY INFORMATION:

Utility

JP 1997-237273 19970902

DOCUMENT TYPE: FILE SEGMENT:

Granted

PRIMARY EXAMINER:

Saunders, David

ASSISTANT EXAMINER:

Tung, Mary Beth

LEGAL REPRESENTATIVE:

Evenson, McKeown Edwards & Lenahan P.L.L.C.

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

26 Drawing Figure(s); 11 Drawing Page(s)

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The methods and compositions of the invention and the compounds used in the invention involve a novel immunosuppression mechanism, accelerated lymphocyte homing immunosuppression (ALH-immunosuppression). For example, the compound FTY720 specifically directs lymphocytes to the peripheral lymph nodes, mesenteric lymph nodes, and Peyer's patches. By reversibly sequestering lymphocytes in these tissues, the compounds can inhibit an immune response in a mammal. Understanding these mechanisms provides a novel immunosuppression therapy that can synergistically interact with other immunosuppressive compounds. Screening methods for identifying similar ALH-immunosuppression compounds are also described. The invention allows better treatments and therapies wherever an immunosuppression regimen is desired.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 19 OF 21 USPATFULL

ACCESSION NUMBER:

1999:106496 USPATFULL

TITLE:

Benzene compound and pharmaceutical use thereof

INVENTOR(S): Fujita, Tetsuro, Muko, Japan

Adachi, Kunitomo, Chikujo-gun, Japan

Kohara, Toshiyuki, Iruma, Japan Kiuchi, Masatoshi, Iruma, Japan Chiba, Kenji, Chikujo-gun, Japan

Teshima, Koji, Iruma, Japan

Mishina, Tadashi, Chikujo-gun, Japan

Yoshitomi Pharmaceutical Industries, Ltd., Osaka, Japan

(non-U.S. corporation)

APPLICATION INFO.: US 1997-801390 19970220 (8)
RELATED APPLN. INFO.: Continuation-in-part of Ser. No. WO 1995-JP1654, filed

on 22 Aug 1995

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PATENT ASSIGNEE(S):

PRIMARY EXAMINER: Raymond, Richard L.

LEGAL REPRESENTATIVE: Evenson, McKeown Edwards & Lenahan P.L.L.C.

NUMBER OF CLAIMS: 37 EXEMPLARY CLAIM: 1 LINE COUNT: 10327

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A benzene compound of the formula ##STR1## wherein each symbol is as defined in the specification; an optically active isomer or salt thereof, a medicinal composition containing the same, and an immunosuppressant containing the same as the active ingredient.

The compound, optically active isomer or salt has an excellent immunosuppressive effect and is useful as an inhibitor for the rejection reaction occurring in organ or bone marrow transplantation, and as a preventive or remedy for articular rheumatism, atopic eczema (dermatitis), Beh.cedilla.et's disease, uveal disease, systemic lupus erythematosus, Sjogren's syndrome, multiple sclerosis, myasthenia gravis, type I diabetes, endocrine ophthalmopathy, primary biliary, cirrhosis, Crohn's disease, glomerulonephritis, sarcoidosis, psoriasis, pemphigus, aplastic anemia, idiopathic thrombocytopenic purpura, allergy, polyarteritis nodosa, progressive systemic sclerosis, mixed connective-tissue disease, aortitis syndrome, polymyositis, dermatomyositis, Wegener's granuloma, ulcerative colitis, active chronic hepatitis, autoimmune hemolytic anemia, Evans' syndrome, bronchial asthma and pollinosis. It is useful also as an antifungal agent and hair growth stimulant.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 20 OF 21 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 2000:527310 BIOSIS DOCUMENT NUMBER: PREV200000527310

TITLE: Recurrent renal allograft rejection: Therapeutic options.

AUTHOR(S): Hauser, Ingeborg A. (1)

CORPORATE SOURCE: (1) Funktionsbereich Nephrologie, Johann Wolfgang

Goethe-Universitaet, Frankfurt/Main Germany

SOURCE: Kidney & Blood Pressure Research, (1999) Vol. 22, No. 4-6,

pp. 259-263. print.

Meeting Info.: Joint Scientific Meeting of the Society for

Nephrology and the German Working Group for Clinical Nephrology Freiburg, Germany September 18-21, 1999

ISSN: 1420-4096.

DOCUMENT TYPE: Conference LANGUAGE: English

SUMMARY LANGUAGE: English ANSWER 21 OF 21 EMBASE COPYRIGHT 2002 ELSEVIER SCI. B.V. 1999049708 EMBASE ACCESSION NUMBER: TITLE: Immunology - Tenth International Congress. AUTHOR: Vohora S.B.; Raisuddin S. CORPORATE SOURCE: S.B. Vohora, Dept. of Med. Elementol./Toxicology, Jamia Hamdard (Hamdard University), New Delhi 110 062, India. root@hamduni.ren.nic.in SOURCE: IDrugs, (1999) 2/1 (22-25). ISSN: 1369-7056 CODEN: IDRUFN COUNTRY: United Kingdom DOCUMENT TYPE: Journal; Conference Article FILE SEGMENT: 026 Immunology, Serology and Transplantation 030 Pharmacology 037 Drug Literature Index LANGUAGE: English SUMMARY LANGUAGE: English This report provides selective coverage of this meeting. Over 2000 participants (including some Nobel laureates) from 67 countries attended. Among these was a significant number from Eastern European countries. This report focuses primarily on DNA vaccines and conventional vaccine development, as well as drug development. It also covers some of the plenary lectures that were delivered by immunology luminaries. The overall focus of the meeting centered on developments in the areas of vaccines, HIV and immune mechanisms. A significant number of presentations concentrated on tumor immunology and immunotherapy. Immunomodulation was another area of major discussion. With such a large, well-attended meeting, it is difficult to provide coverage for each of the speakers in a particular section, and the omissions are unintentional. => d kwic 20-21 ANSWER 20 OF 21 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. & Systems of Organisms renal artery: circulatory system, excretory system, structural changes CMV infection [cytomegalovirus infection]: concomitant disease, viral disease; acute rejection: immune system disease, pathogenesis, urologic disease; chronic rejection: immune system disease, pathogenesis, urologic disease; late renal allograft loss: immune system disease, prevention, urologic disease; polyoma virus infection: concomitant disease, viral disease; recurrent renal allograft rejection: immune system disease, pathogenesis, treatment options, treatment outcomes, urologic disease; renal allograft rejection: classifications, diagnosis,. RN 104987-11-3 (FK506) 104987-11-3 (TACROLIMUS) 162359-56-0 (FTY720) 59865-13-3 (CYCLOSPORINE A) 216973-42-1 (CSA) 1247-42-3 (METHYLPREDNISONE) 128794-94-5 (MYCOPHENOLATE MOFETIL) 53123-88-9 (RAPAMYCIN) 53123-88-9 (SIROLIMUS) ANSWER 21 OF 21 EMBASE COPYRIGHT 2002 ELSEVIER SCI. B.V.

CT

Medical Descriptors:

*immunology
*drug research

human immunodeficiency virus 1

tumor immunology immunomodulation vaccination drug manufacture delayed hypersensitivity drug design immune response cross reaction cytokine production human clinical trial meta analysis multicenter study human tissue human cell conference paper *DNA vaccine: CT, clinical. .

RN (2 amino 2 [2 (4 octylphenyl)ethyl] 1,3 propanediol) 162359-56-0; (gonadorelin) 33515-09-2, 9034-40-6; (interleukin 2) 85898-30-2; (milodistim) 137463-76-4; (lymphotactin) 156561-05-6; (thalidomide derivative) 31804-66-7

CN (1) Fty 720; (2) Cc 3052

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	ENTRY	SESSION
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